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NOVEL SEPARATION MATRIX

Technical field

The present invention relates to a novel separation matrix for isolation of target molecules, such as proteins, from a liquid. The invention also encompasses a method of preparing such a matrix as well as a chromatography column packed with such as separation matrix.

Background

Biotechnological methods are used to an increasing extent in the production of proteins, peptides, nucleic acids and other biological compounds, for research purposes as well as in order to prepare novel kinds of drugs. Due to its versatility and sensitivity to the compounds, chromatography is often the preferred purification method in this context. The term chromatography embraces a family of closely related separation methods, which are all based on the principle that two mutually immiscible phases are brought into contact. More specifically, the target compound is introduced into a mobile phase, which is contacted with a stationary phase. The target compound will then undergo a series of interactions between the stationary and mobile phases as it is being carried through the system by the mobile phase. The interactions exploit differences in the physical or chemical properties of the components in the sample.

In liquid chromatography, the target compound is present in a liquid together with one or more contaminants or undesired substances. Said liquid is contacted with a stationary phase, known as a matrix, which is commonly comprised of either a collection of homogenous, porous or non-porous particles or a monolith of organic or inorganic origin. The properties of the separation matrix will in large decide the efficiency obtained when used in a separation process, such as chromatography. Usually, a separation matrix is comprised of a support to which groups capable of interaction with the target and known as ligands have been coupled. Thus, the ligands will impart to the supports the ability to

effect the separation, identification, and/or purification of molecules of interest. In the prior art, a number of different techniques for controlling the density of ligands on a support have been suggested, which techniques generally fall into one of the following four categories:

a) Manipulation of reaction conditions which activate the matrix, i.e. which introduce a reactive group which can couple the ligand. This often involves varying the concentration of activating reagents, reaction time, reaction temperature, pH, or combinations of these variables. Thus, the efficiency of the reaction, i.e. the extent of desired reaction as opposed to competing side reactions, will be strongly influenced by reaction conditions.

b) Manipulation of reaction conditions during the actual coupling of the ligand to the support. This may involve varying the concentration and/or the total amount of ligand the support is challenged with, ionic strength of the coupling buffer, and type of salt in the coupling buffer as well as the variables of time, temperature, pH, etc., mentioned above. Similarly to the technology described above, due to the strong influence of the reaction conditions, this method may also prove difficult to apply in a practical and reproducible manner.

c) Manipulation of the amount of reactive or activatable groups incorporated into the support by varying composition at the time of its formation. For a polymeric support, this would include varying the nature and/or amount of monomer during the polymerisation. Obviously, one should subsequently apply the techniques of a and/or b above in a second step to couple the ligand.

d) For polymeric ligands, manipulation of the amount of ligand incorporated into the polymer by preparation of a polymerisable ligand monomer and varying the concentration of this monomer in the monomer feed during polymerisation. A drawback with this technology is that many ligands useful for chromatographic separations contain functional groups which are incompatible with

the conditions necessary for formation of the desired polymer, such as by being unstable under the contemplated polymerisation conditions or by interfering with the polymerisation reaction.

A different method of controlling the density of ligands on a support was suggested in US 5,561,097 (Gleason et al.), which relates to a method of providing an optimised ligand density on a polymeric support, which method is stated to be obtained in a practical and reproducible manner. This can be achieved by a method comprising a step of reacting ligand and a quencher molecule with activated sites on an azlactone functional support under conditions that promote competition of ligand with quencher for the activated sites. One advantage presented is that the method is a single step procedure, without any need of a separate step to activate or deactivate reaction sites on the support. The method disclosed is stated to be especially advantageous for the coupling of small molecules. A disadvantage of this method is that in order not to favour ligand over quencher, an understanding of the reaction kinetics for ligand and reaction kinetics for quencher will be required, including the rate constant for coupling, the concentration of ligand, the nucleophilicity of ligand and quencher, etc.

Further, in biotechnological preparation of target molecules such as proteins, it is well known that to enable an efficient purification thereof, a series of two or more process steps utilising different kinds of separation matrices is often required. US 6,426,315 (Bergström et al.) suggests to replace such a series of steps by using a multifunctional porous separation matrix, i.e. to present the different kind of matrices on a single separation matrix. More specifically, US 6,426,315 relates to a process for preparing such multifunctional porous separation matrices by introducing different functionalities in different layers of the matrix. In brief, the process includes to contact a separation matrix that comprises reactive groups with a reagent, the amount of which is not sufficient for reaction with all groups present in the matrix, and wherein the reaction between the reagent and said reactive groups is rapid compared to the mass transport of

the reagent within the matrix. The reactive groups may e.g. be hydroxyl groups, double bonds etc, while the reagent may be a compound that introduces a desired functionality within the matrix, directly or indirectly. In the last mentioned case, the reagent is a compound known as an activating agent, such as a halogenating agent, and the desired functionality is then introduced in a subsequent step. The most preferred functionalities are groups that provide desired separation characteristics to the matrix, commonly known as ligands. Alternatively, the functionalities introduced are the degree of crosslinking, the density or the porosity of the matrix. In order to provide further layers, the reactive groups may be further reacted with another reagent. Thus, the method according to US 6,426,315 may exhibit the drawbacks discussed above under a) and b). In addition, even though the teachings of US 6,426,315 enable the construction of a separation matrix which exhibits a multitude of functions, each one of which will provide different properties as regards binding and diffusion in a separation process, there is no guidance in US 6,426,315 with regard to how to manufacture a separation matrix to that provides an optimal mass transport within the matrix. Thus, there is still a need in this field of alternative methods of producing separation matrices with improved such properties.

In US 5,945,520 (Burton et al.), it is stated that a problem with the known kind of multi or mixed mode chromatography matrices that adsorb a target compound via hydrophobic interactions is that binding efficiencies of less hydrophobic targets will be low unless high salt concentrations are used. To avoid such necessary addition of salt, US 5,945,520 suggests a chromatographic resin, which presents an ionisable ligand comprised of an ionisable functionality and a spacer arm, which attaches said functionality to a solid support matrix. The ionisable functionality is partially electrostatically charged at the pH of adsorption of the target compound to the resin, and is either further charged or of opposite charge at the pH of desorption of the compound from the resin. The ionisable functionalities are selected from a specified group of possible functionalities. In one embodiment, the ionisable functional group is derived

from either 2-mercapto-1-methylimidazole or (-)phenylpropanolamine and coupled to a density of at least 150 µmol per millilitre of resin. Such a high ligand density is stated to provide a sufficient hydrophobicity to adsorb target compounds without the need of adding excessive amounts of salt to the liquid. Thus, US 5,945,520 discloses multifunctional ligands evenly coupled to a resin, which consequently can be described as a homogenous separation matrix.

Accordingly, there is still a need in this field of novel separation matrices, which avoid one or more of the problems associated with the prior art. For practical and economical reasons, there is a need in this field of separation matrices that provides an improved mass transport of target molecules.

Summary of the invention

One object of the present invention is to provide a separation matrix, wherein the mass transport properties during adsorption and/or desorption have been improved. This can be achieved by a separation matrix as described in claim 1. Thus, an object of the invention is to provide a separation matrix, which is especially advantageous for use in large-scale purification of a target molecule.
A specific object of the present invention is to provide a separation matrix comprised of essentially spherical particles, wherein the mass transport properties differ along the radius of each such particle.
Another object of the present invention is a separation matrix, which allows elution of target molecules at an increased efficiency.

A further object of the invention is to provide a method of preparing a separation matrix comprised of a porous support, wherein a ligand density gradient is provided. This can be achieved by careful control of the reactivity of ligand during coupling thereof to the support.
Yet a further object of the invention is to provide a process of liquid chromatography, wherein the mass transport during adsorption and/or desorption is improved.

Thus, one object of the invention is to provide a process of liquid chromatography, wherein the mass transport during adsorption and/or desorption is controllable.

Further objects and advantages of the present invention will appear from the detailed description that follows.

Brief description of the drawings

Figure 1 shows in a schematic way how the adsorption of a target molecule can be facilitated by an increasing ligand density towards the centre of a particulate separation matrix.

Figure 2A-H show various illustrative chemical gradients in that can be provided in essentially spherical particles according to the invention. Two ligand kinds are shown, ligand 1 (line) and ligand 2 (broken line). The gradients are shown in diagrams having Relative density on the Y-axis and a dimensionless radius on the X-axis.

Definitions

The term "gradient" means a change in the value of a quantity with change in a given variable, especially per unit distance, in a specified direction.

The term a "chemical gradient" means herein that the chemical properties vary in a systematic manner.

The term a "continuous and smooth" gradient means that it increases or decreases in a continuous fashion; linearly or with varying degrees of convexity or concavity. Thus, a "continuous and smooth" gradient is essentially free from discontinuous steps. In the context of essentially spherical particles, a "radial" gradient means that the gradient increases or decreases towards the centre of the beads.

The term "ligand" means a chemical entity, which comprises at least one functional group capable of interaction with a target molecule. In this context, it is

understood that a "ligand" may or may not comprise a spacing element, which distances the functional groups from the surface of a solid support.
The term "affinity ligand" means herein a ligand comprising a functionality capable of interaction via biological affinity with a target molecule, such as biotin interacting with streptavidin, an antibody interacting with an antigen etc.
The term "ligand density" means herein the degree of substitution of a support, such as a particle, and is commonly measured in µmol/ml resin.
The "surface" of a porous support refers to the outer surface as well as to the pore surfaces.

Detailed description of the invention

In a first aspect, the present invention relates to a separation matrix comprised of ligands coupled to the surfaces of a porous support, wherein the ligands provide at least one chemical gradient in the support, which gradient may be directed across the support or be present within the support, depending on its nature. Illustrative supports are e.g. membranes, monoliths and particles. Thus, in an advantageous embodiment, the present invention is a separation matrix comprised of ligands coupled to the surfaces of at least one porous particle, wherein the ligands provide at least one chemical gradient between the centre and the exterior surface of each porous particle. In this context, it is understood that a chemical gradient means a systematic and repeated change in any chemical property, preferably a property that is utilised in, or influence, a subsequent separation process. Illustrative chemical properties are concentration, density and attraction. The chemical gradient present in the particles according to the invention is preferably created in a way that improves the mass transport when the matrix is utilised for separation, such as in chromatography. Accordingly, even though heterogeneous particles may have been resulted from prior art methods of manufacture, the present invention suggests for the first time heterogeneous particles which have been specifically designed to incorporate a chemical gradient that alters and preferably improves their adsorption and/or desorption properties during a separation process. Illustrative adsorption and/or

desorption properties are e.g. capacity, selectivity, efficiency, intraparticle mass transport, rigidity etc.

The separation matrix according to the present invention is useful in separation and/or purification of practically any target molecule or compound, such as in chromatography or membrane separation. Illustrative examples of target molecules are biomolecules, such as proteins, peptides, nucleic acids, plasmids and virus, as well as organic molecules. A process of liquid chromatography utilising the present separation matrix will be discussed in more detail below.

In the most advantageous embodiment of the present separation matrix, the porous particle is an essentially spherical bead. Such beads may be made of an inorganic material, such as silica, or of an organic material, and are preferably natural or synthetic polymers.

Thus, in one embodiment, the beads are comprised of a cross-linked carbohydrate material, such as agarose, agar, cellulose, dextran, chitosan, konjac, carrageenan, gellan, alginate etc. Such beads are easily prepared according to standard methods, such as inverse suspension gelation (S Hjertén: Biochim Biophys Acta 79(2), 393-398 (1964). Alternatively, the beads according to the invention are made from commercially available products, such as Sepharose™ FF (Amersham Biosciences AB, Uppsala, Sweden).

In another embodiment, the polymer beads are comprised of cross-linked synthetic polymers, such as styrene or styrene derivatives, divinylbenzene, acrylamides, acrylate esters, methacrylate esters, vinyl esters, vinyl amides etc. Such polymers are easily produced according to standard methods, see e.g. "Styrene based polymer supports developed by suspension polymerization" (R Arshady: Chimica e L'Industria 70(9), 70-75 (1988)). Alternatively, a commercially available product, such as Source™ or Sephacryl™ (Amersham Biosciences AB, Uppsala, Sweden) can be utilised as starting material to prepare the beads according to the invention.

In an advantageous embodiment of the present separation matrix, at least one chemical gradient is a ligand density gradient resulting from a varying density of the ligands present on the support, such as on each porous particle.
In an alternative embodiment, at least one chemical gradient is the result of varying pKa values of the functional groups of the ligands present on each porous particle. In yet an alternative embodiment, at least one chemical gradient is the result of a varying net charge of the ligands present on each porous particle. Such gradients can be prepared in methods analogue to the ones described in relation to ligand density gradients, wherein the pKa values and net charge are controlled instead of the density of coupled ligands.

A method of preparing chemical gradients according to the invention in a porous support will be described in detail below.

The chemical gradient may be of either direction, i.e. directed and hence increasing towards the centre of the support, or directed towards the outer surface of the support and consequently decreasing towards the centre. As the skilled person in this field will realise, as the mass transport is increased towards the centre of a support, the increasing amount of target molecule in the inner will give rise to a concentration gradient, which then works against the desired direction of mass transport. However, in most cases, such a concentration driven force will be smaller than the driving force into the desired direction. If necessary, the skilled person in this field can take this into consideration when the support is designed and ensure that the attraction force provided by the functional groups is sufficient. In the case of an opposite gradient, such a concentration gradient may instead cooperate with the driving force of the mass transport.

Thus, in an illustrative embodiment of the present separation matrix, a ligand density gradient is designed in porous essentially spherical particles to maximise the mass transport in the outermost part, corresponding to about 40% of

the radius. This means that the ligand density increases towards the centre and the slope of the ligand gradient is largest in the outer part of the particles and lowest in the inner part of the particles. The outermost part corresponding to the outer 40% of the particle radius accommodates almost 80% of the total particle volume and protein capacity, respectively. This means that it will not be as important to utilise the capacity in the inner part of the particles. Therefore, the above suggested gradient can also be designed in a way that the slope of the ligand density gradient increases from the outer surface of the particle to about 40% of the radius and then the ligand density will decrease towards zero in the centre of the particles. In a specific embodiment of the present separation matrix, the support is comprised of essentially spherical particles, wherein each particle is comprised of a solid non-porous material surrounded by a porous material as discussed above and wherein at least one chemical gradient has been provided in the porous part. This embodiment may advantageously be used in applications where a high flow rate is desired, since a solid inner part will improve the rigidity of the particle while an efficient separation can be obtained by using only a part of the particle, such as about 40% of the radius, as explained above.

In the most advantageous embodiment of the present separation matrix, at least one chemical gradient is a continuous and smooth gradient. Such gradient(s) may be present only in the outer part of a support, as discussed above. Alternatively, such gradient(s) may not be present only in the inner part of a particle, such as leaving an outer shell of a porous particle wherein there is no gradient. As will be discussed in more detail below, the present separation matrix may comprise two or more chemical gradients, such as one or more continuous and smooth gradients and one or more gradients that comprise discontinuous steps. The skilled person in this field will be able to decide the most advantageous extent of the chemical gradient(s) for each intended purpose, i.e. depending on which separation property it is most desired to alter or improve.

In one embodiment of the present separation matrix, the ligands of each particle provide at least two different functionalities. In one embodiment, said functionalities are selected from the group that consists of cation exchange ligands, anion exchange ligands, hydrophobic interaction chromatography (HIC) ligands, reversed phase chromatography (RPC) ligands, immobilised metal chelating ligands (IMAC), thiophilic ligands, and affinity ligands. Such functionalities are well known to the skilled person in this field and are easily prepared by standard methods, see e.g. Janson and Rydén in Protein Purification: Principles, High Resolution Methods, and Applications (1989 VCH Publishers, Inc).

In a specific embodiment of the present separation matrix, said at least two different functionalities are present on the same ligand. Thus, the ligands may comprise groups that are partly electrostatically charged, or partly electrostatically charged, at the pH of binding of the target molecule to the matrix and either further charged or of an opposite charge at the pH of elution. In an illustrative embodiment, the ligands present zwitterionic functionalities. Thus, in this embodiment, the ligand will comprise two groups that are cationic and anionic, respectively, at a specified pH value. Consequently, the adsorption may then utilise ionic interactions of a first charge, while elution can be provided by a change in pH to provide desorption of the target molecule from the matrix by neutralisation of said first charge and repulsion by ionic interactions of the second charge, which is opposite from the first. Examples of zwitterions are numerous and well known to the skilled person in the art, such as the common amino acids, and may be exemplified e.g. by N-[tris-(hydroxymethyl)methyl]-3-aminopropane sulphonic acid (TAPS), dimethyl glycin and glycin alanine.

In an alternative specific embodiment, said at least two different functionalities are present on different ligand kinds, and each such ligand kind provides a separate chemical gradient within the porous particle. Accordingly, in an illustrative embodiment, the present separation matrix comprises two or more

chemical gradients provided by two or more different kind of ligands, such as two ligand density gradients of the same or opposed direction, wherein each ligand kind provides a separate chemical gradient.

As the person skilled in the art will understand, the discussion below will apply equally well to the embodiment where a chemical gradient has been obtained from ligands comprising more than one functionality and to the embodiment using different kinds of ligands. Thus, by combining two different chemical gradients, one for a positive group and the other for a negative group, the present invention allows to increase the mass transport of a target molecule, such as a charged protein, in the outer part (corresponding to about 40% of the radius of an essentially spherical particle) and then prevent the protein to be transported into the centre of the support by charge repulsion.

If positively charged proteins are the target molecules, the gradient of negatively charged ligands is constructed as described above. The slope of the negatively charged ligand gradient increases up to a value corresponding to about 40% of the radius of an essentially spherical particle, and then the ligand density decreases towards zero in the centre of the support. This means that the ligand density is at a maximum value at a value corresponding to about 40% of the radius of an essentially spherical particle. The positively charged ligand gradient may begin at a location corresponding to about 40% of the radius of an essentially spherical particle and increases toward the centre of the support. This gradient is designed in a way so that the positively charged proteins are hindered by charge repulsion to penetrate the centre of the support.

In a second aspect, the present invention relates to a chromatography column packed with a separation matrix comprised of ligands coupled to the surfaces of a porous support, wherein the ligands provide at least one chemical gradient within the support.

In an advantageous embodiment, at least one chemical gradient is a ligand density gradient. In one embodiment, the present chromatography column has been packed with a separation material as described above. The chromatography columns according to the invention may be relatively small and useful for laboratory applications or of a larger size suitable for production of target molecules in large scale. The column material and design may be adapted depending on the intended application using standard materials and technologies.

In a third aspect, the present invention relates to a method of preparing a separation matrix that comprises ligands coupled to the surfaces of a porous support, which method comprises the step of

(a) providing activatable groups on the surface of a porous support;

(b) activating said groups with an activation agent;

(c) reacting groups activated according to step b) with a compound which comprises at least one functionality;

wherein control of the reactivity in step (c) results in at least one chemical gradient within the support. Alternatively or additionally, the gradient(s) are obtained by controlling the diffusion rate during step (c). The control of step (c) can also be defined as control of the reaction rate. The skilled person in this field can adjust the appropriate parameters to obtain a suitable relationship between reactivity and diffusion. In one embodiment, the reactivity is slightly higher than the diffusion into the support.

The gradients may be generated by using a limiting amount of compound comprising functionalities (ligand) in step (c) as compared to the activated groups resulting from step (b). It is also possible to control other parameters such as the temperature, the concentration of the ligand or the type of solvent to control the diffusion and/or the reactivity of the ligand, creating gradients with different ligand substitution profiles. Using different limiting quantity of ligand can also generates different of gradient profiles. Thus, in one embodiment, the diffusion rate is controlled by adjustment of the concentration of ligand.

The porous support may be as discussed above in relation to the first aspect of the invention. In an advantageous embodiment, the support is comprised of at least one porous particle, preferably an essentially spherical particle, and at least one chemical gradient extends between the centre and the exterior surface of each porous particle. Various general techniques for preparing a porous particle that carries ligands are well known in this field. As is easily realised, the method of choice will depend on the nature of the porous particle.

The activatable groups provided in step (a) may be any groups commonly used in this context, such as carboxylic groups (activatable with NHS/EDC), amines, allyl groups etc. Thus, in an advantageous embodiment of the present method, the activatable groups of step (a) are carbon-carbon double bonds. Thus, the porous support may for example be particles made from a synthetic polymer, wherein double bonds such as vinyl groups left unreacted in the preparation thereof are easily available.

In one embodiment, the present method also comprises to provide the activatable groups present at the surface of at least one porous particle in a step preceding step (a). The allylation is easily performed following standard methods with a suitable agent, such as with allyl glycidyl ether (AGE) or allyl bromide. This may for example include to allylate suitable groups such as hydroxyl groups, which are available in most natural polymers such as polysaccharides, e.g. agarose. However, other activatable groups are well known to the skilled person in this field, and illustrative further examples are e.g. amines, thiols, carboxy groups etc. Accordingly, in one embodiment, the step preceding step (a) comprises to allylate hydroxyl groups present on the surface of a porous support.

In an alternative embodiment, steps (a)-(c) above are replaced by a single step, wherein an activated ligand is reacted with the activatable groups present on

the surface of the support. This may for example involve use of a compound, which carries both a reactive group, such as an epoxide, which is available for reaction with the e.g. a hydroxyl group on the surface of the support, and one or more ligand functionalities.

Step (b) can be performed using any commonly used activation agent capable of rendering a carbon-carbon double bond reactive. In one embodiment, the activation agent is a free radical. In another embodiment, the activation agent comprises an electrophile, such as a halogen, e.g. bromine, chlorine or iodine, or a hydroxide thereof. Thus, in one embodiment, the activation agent used in step (b) is a halogen. Activation of a carbon-carbon double bond with a halogen or halogenating agent may result in halohydrines, which are easily converted to reactive epoxy groups in accordance with standard methods. Thus, in a specific embodiment, step (b) also includes to convert the halogenated product to an epoxy group.

As the skilled person in this field will easily realise, in addition to the actual functional group(s), the compound coupled in step (c) may also comprise an element which distances said functional group(s) from the support surface. Such elements are commonly used and known as spacer elements or spacers. Alternatively, a spacer may coupled to the surface of the particle surface before the activation thereof. As is well known, any such distancing element will work as a spacer, regardless of the method for its introduction, and in principle any group or compound that provides such distancing will be denoted a spacer. As appears from the above, some of the more frequently used activating schemes comprise an activating agent such as allyl glycidyl ether (AGE), which in part will be transformed into a spacer.

In one embodiment, at least one chemical gradient is a continuous and smooth ligand density gradient, as discussed above in the context of the first aspect of the invention.

In the most advantageous embodiment of the present method, at least one chemical gradient is a ligand density gradient. The present method may be repeated in order to obtain two or more chemical gradients, as discussed above. In an alternative embodiment, the gradients are obtained more or less simultaneous, in a single process.

In one embodiment of the present method, the functionalities of in step (c) provide at least two different functionalities. Such functionalities may be as described above in the context of the first aspect of the invention. In one embodiment, said at least two different functionalities are present on the same ligand, such as on a zwitterionic compound.

In an alternative embodiment, said at least two different functionalities are present on different ligand kinds, and a separate chemical gradient is provided for each such ligand kind within the support. Such different gradients may be as discussed above in the context of the first aspect of the invention. Thus, in one embodiment, two chemical gradients are provided in the support, one of which is a ligand density gradient. The effects and advantages of such gradients and combinations of gradients will be discussed in more detail below in the context of the process of liquid chromatography.

The present invention also encompasses a separation matrix prepared using the present method.

In a fourth aspect, the present invention relates to a process of liquid chromatography, wherein a liquid comprising a target molecule is contacted with a separation matrix that comprises ligands coupled to the surfaces of a porous support and the ligands provide a chemical gradient within the support. In one embodiment, the support is comprised of at least one porous particle, and a chemical gradient extends between the centre and the exterior surface of each

porous particle. Thus, the gradient may be a radial gradient. The nature of the separation matrix may be as described above in the context of the first aspect of the invention. The general principles behind liquid chromatography are well known to the skilled person in this field and described in various textbooks.

In the most advantageous embodiment, the present process further comprises a step of eluting the adsorbed target molecule from the matrix by contacting the matrix with an eluent. The nature of the eluent will obviously depend upon the nature of the ligands coupled to the particles. For example, in case of ionexchange gradient elution, the eluent may comprise a salt gradient or a pH gradient. In a specific embodiment of the present process, the functional groups of the ligands are zwitterions and the elution is performed at a pH different from that used during the adsorption.

In one embodiment, in the separation matrix used in the present process, the ligand density increases towards the centre of the support. Hence, if the particles are essentially spherical beads, this embodiment includes a radial gradient. In an alternative embodiment, the gradient is of the opposite direction. As described above in the context of the first aspect of the invention, the present separation matrix may contain two or more chemical gradients, one of which is preferably a continuous and smooth gradient.

By utilising a separation matrix comprising two different ligand gradients, where each of the two ligands have different chromatographic properties, both the adsorption and the desorption processes can be improved. For example, by choosing one ligand that attracts and one ligand that repels the target molecules, the adsorption can be directed to the outer part of the support and the target molecules are prevented from penetrating the inner part of the support. The gradient of the attracting ligand is adjusted to increase the mass transport of the sample molecules in the outer part of the support.

Another advantage of using two ligands and two different gradients is the possibility to achieve a separation of target molecules in each support, e.g. in each particle, and in this way increase the selectivity of the chromatographic process, such as a separation in a column. This can, for example, be accomplished by combining one gradient of an ion-exchange functionality and another gradient of a functionality that supports hydrophobic interactions, i.e. a HIC ligand. The gradient of the HIC-ligands decreases towards the centre of the particles and the gradient of the ion-exchange ligand increases towards the centre. This means that hydrophilic target molecules, which are oppositely charged to the ion-exchange group, will mainly be adsorbed in the inner part of the particles and the hydrophobic charged sample molecules will be adsorbed in the outer part. When such particles according to the invention are used in column liquid chromatography, the adsorbed target molecules are easily desorbed by gradient elution. To fully take advantage of the protein separation in the particles, the solvent gradient should be designed in a way that the proteins adsorbed in the outer part of the particles are eluted first and the proteins adsorbed in the inner part of the particles are eluted last.

As mentioned above, the present invention is useful for separation of virtually any target molecule, depending on how the separation matrix is designed. In one embodiment, the target molecule is a protein. The present process of liquid chromatography may either be a process of purifying one desired target molecule from a solution, such as a fermentation broth, or a process used for purification of a liquid by removal of an undesired target molecule, such as a virus.

Detailed description of the drawings

Figure 1 shows in a schematic way how the adsorption of a target molecule can be facilitated by an increasing ligand density towards the centre of a particulate separation matrix. The drawing is intended to illustrate an embodiment where the functional groups of the ligands are surrounded by a field, such as the field

surrounding a charged functionality. The arrows indicate how an increasing force will facilitate the mass transport into the particle.

Figure 2A-H shows illustrative examples of chemical gradients provided according to the present invention. Figure 2A shows a gradient which increases towards the centre of the support, which gradient starts within the support and hence leaves an outer part of the particle without ligands; Figure 2B shows a gradient which decreases towards the centre of the support; Figure 2C shows a gradient which reaches a peak value halfway to the centre of the support; Figure 2D shows a gradient which decreases towards the centre of the support, i.e. where the ligands are present at a higher concentration in the outer part of the support; Figure 2E shows two gradients of opposite direction within the support; Figure 2F shows two gradients of the same direction but of different inclination within the support; Figure 2G illustrates two gradients, one is in the centre of the particle and one surrounding the centre, similar to Figure 2C; and Figure 2H shows a support, wherein one ligand or functionality describes a planar curve and the other is a gradient similar to the one described in Figure 2B. A support having the kind of gradients shown in Figure 2H can e.g. be obtained by starting from a support material which has already been functionalised with ligand to a constant substitution degree and subsequently applying the method according to the invention to provide a gradient.

EXPERIMENTAL PART

The present examples are provided for illustrative purposes only, and shall not be construed as limiting the invention as defined by the appended claims. All references given below and elsewhere in the present application are hereby included herein by reference.

Linear gradient in bead

General:

The volumes of matrix refer to settled bed volume. The weights of matrix given in gram refer to suction (water pump) dry weight. It is understood that these matrices are still water solvated material.

Stirring is referring to a suspended, motor-driven stirrer since the use of magnet bar stirrer is prompt to damage the beads.

Analysis of the functionality and the determination of the degree of allylation, epoxidation, or the degree of substitution of ion exchanger groups on the beads refer to conventional methods.

One way to prepare some bead with increasing or decreasing linear gradient from the surface to the inside of the bead is exemplified below with a Sepharose™ 6 FF gel (Amersham Biosciences, Uppsala, Sweden).

A specific example is described for anion and cation exchange media.

The gradients were generated by using a limiting amount of ligand as compared to the activated allyl groups. It is also possible to take advantage of other parameters such as the temperature, the concentration of the ligand or the type of solvent to control the diffusion and the reactivity of the ligand, creating gradients with different ligand substitution profiles.

Using different limiting quantity of ligand can also generates different of gradient profiles.

Example 1. Introduction of allyl group on the matrix:

In a typical procedure allylation was carried out with allyl glycidyl ether, but note that the introduction of allyl groups on the solid support can as well be easily achieved with using allyl bromide.

Activation of Sepharose™ with allyl glycidyl ether:

A 80 g quantity of Sepharose™ 6 FF was mixed with 0.5 g of $NaBH_4$, 13 g of Na_2SO_4 and 40 ml of 50% aqueous solution of NaOH. The mixture was stirred for 1 hour at 50 °C. After addition of 100 ml of allylglycidyl ether the suspension was left at 50 °C under vigorous stirring for an additional 18 hours. After filtration of the mixture, the gel was washed successively, with 500 ml distilled water, 500 ml ethanol, 200 ml distilled water 200 ml 0.2 M acetic acid and, 500 ml distilled water.

Titration gave a degree of substitution of 0.3 mmol of allyl/ml of gel.

Note: By adding different quantities of NaOH, the degree of substitution can be varied.

For example: With exactly the same method as above the use of 50 ml of 50% NaOH will give a degree of substitution of around 0.4 mmol of allyl/ml of gel.

Example 2. Linear gradient with decreasing ligand substitution degree towards the middle of the bead.

Example 2.1. Cation exchange media

In a typical procedure the carboxylic groups were introduced on the matrix with thiol containing derivatives but derivatives containing other reactive nucleophilic groups can as well be used.

<u>Gradients of immobilised 3-mercaptopropionic acid:</u>

Bromine was added to a stirred suspension of 100 ml of allyl activated Sepharose™ 6 FF (0.3 mmol of allyl/ml), 4 g of AcONa and 100 ml of distilled water, till a persistent yellow colour was obtained. Sodium formate was then added till the suspension was fully decolourised.

The reaction mixture was filtered and the gel washed with 500 ml of distilled water. The activated gel was then directly transfer to 4 distinct reaction vessels:

A) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 366 µl of thiopropionic acid (0.7 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.
The reaction was left 10 hours under stirring at 70 °C. Filtration of the reaction mixture and washing with 100 ml of distilled water gave the thiopropionic Sepharose derived gel.

B) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 366 µl of thiopropionic acid (0.7 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.
The reaction was left 10 hours under stirring at 30 °C. Filtration of the reaction mixture and washing with 100 ml of distilled water gave the thiopropionic Sepharose derived gel.

C) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 262 µl of thiopropionic acid (0.5 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.
The reaction was left 10 hours under stirring at 70 °C. Filtration of the reaction mixture and washing with 100 ml of distilled water gave the thiopropionic Sepharose derived gel.

D) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 262 µl of thiopropionic acid (0.5 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.
The reaction was left 10 hours under stirring at 30 °C. Filtration of the reaction mixture and washing with 100 ml of distilled water gave the thiopropionic Sepharose derived gel.

Example 2.2. Anion exchange media

In a typical procedure, the anion exchanger groups were introduced on the matrix directly by immobilisation of a tertiary amine, but ligands containing other reactive nucleophilic groups and can as well be used as long as they also present an anion exchanger group or the possibility to generate one.

Gradients of immobilised trimethylamine:

Bromine was added to a stirred suspension of 100 ml of allyl activated Sepharose™ 6 FF (0.3 mmol of allyl/ml), 4 g of AcONa and 100 ml of distilled water, till a persistent yellow colour was obtained. Sodium formate was then added till the suspension was fully decolourised.

The reaction mixture was filtered and the gel washed with 500 ml of distilled water. The activated gel was then directly transfer to 4 distinct reaction vessels:

A) 20 ml of drained brominated gel was charged in to a 100 ml three necked round flask provided with a propeller stirrer together with 8 ml water. A 50/50 solution of sodium hydroxide in water, prepared from 14.9 g NaOH and 0.01g NaBH4 in 14.9 g water, was added drop by drop. An aqueous solution (2 ml) of 0.4 g trimethyl ammonium chloride (0.7 equivalents per allyl group) was then added.

Reaction was run under stirring at 50 °C for 5 hours.

The reaction was terminated by washing the gel on a glass filter funnel with 400 mL water, 80 ml 1M Sodium chloride and another 400 mL of water.

B) 20 ml of drained brominated gel was charged in to a 100 ml three necked round flask provided with a propeller stirrer together with 8 ml water and a 50/50 solution of sodium hydroxide in water, prepared from 14.9 g NaOH and 0.01g NaBH4 in 14.9 g water, was added drop by drop. An aqueous solution (2 ml) of 0.4 g trimethyl ammonium chloride (0.7 equivalents per allyl group) was then added.

Reaction was run under stirring at 20 °C for 5 hours.
The reaction was terminated by washing the gel on a glass filter funnel with 400 mL water, 80 ml 1M Sodium chloride and another 400 mL of water.

C) 20 ml of drained brominated gel was charged in to a 100 ml three necked round flask provided with a propeller stirrer together with 8 ml water and a 50/50 solution of sodium hydroxide in water, prepared from 14.9 g NaOH and 0.01g NaBH4 in 14.9 g water, was added drop by drop. An aqueous solution (2 ml) of 0.286 g trimethyl ammonium chloride (0.5 equivalents per allyl group) was then added.
Reaction was run under stirring at 50 °C for 5 hours.
The reaction was terminated by washing the gel on a glass filter funnel with 400 mL water, 80 ml 1M Sodium chloride and another 400 mL of water.

D) 20 ml of drained brominated gel was charged in to a 100 ml three necked round flask provided with a propeller stirrer together with 8 ml water and a 50/50 solution of sodium hydroxide in water, prepared from 14.9 g NaOH and 0.01g NaBH4 in 14.9 g water, was added drop by drop. An aqueous solution (2 ml) of 0.286 g trimethyl ammonium chloride (0.5 equivalents per allyl group) was then added.
Reaction was run under stirring at 20 °C for 5 hours.
The reaction was terminated by washing the gel on a glass filter funnel with 400 mL water, 80 ml 1M Sodium chloride and another 400 mL of water.

Example 3. Linear gradient with increasing ligand substitution degree towards the middle of the bead.

In a typical procedure, the increasing gradient was realised by first creating a decreasing gradient of a neutral ligand and the resulting gel was further derivatised on the remaining active groups by the chromatographically active ligand.

Example 3.1. Cation exchange media

Gradients of immobilised 3-mercaptopropionic acid and 3-mercapto-1,2-propanediol:

Bromine was added to a stirred suspension of 100 ml of allyl activated Sepharose™ 6 FF (0.3 mmol of allyl/ml), 4 g of AcONa and 100 ml of distilled water, till a persistent yellow colour was obtained. Sodium formate was then added till the suspension was fully decolourised.

The reaction mixture was filtered and the gel washed with 500 ml of distilled water. The activated gel was then directly transfer to 4 distinct reaction vessels:

A) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 251 µl of 3-mercapto-1,2-propanediol (0.5 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.

The reaction was left 1 hour under stirring at 50 °C. After 1 hour the reaction was treated with an aqueous solution (5ml dist.water) of 1.6 ml of thiopropionic acid (3 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition. The reaction was then left 18 hours at 50 °C.

Filtration of the reaction mixture and washing with 100 ml of distilled water gave the thiopropionic Sepharose derived gel.

B) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 251 µl of 3-mercapto-1,2-propanediol (0.5 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.

The reaction was left 1 hour under stirring at 20 °C. After 1 hour the reaction was treated with an aqueous solution (5ml dist.water) of 1.6 ml of thiopropionic acid (3 equivalents per allyl group) and 2.4 g of NaCl which pH was ad-

justed to 11,5 with 50 % aq. NaOH before the addition. The reaction was then left 18 hours at 50 °C.
Filtration of the reaction mixture and washing with 100 ml of distilled water gave the thiopropionic Sepharose derived gel.

C) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 151 µl of 3-mercapto-1,2-propanediol (0.3 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.
The reaction was left 1 hour under stirring at 50 °C. After 1 hour the reaction was treated with an aqueous solution (5ml dist.water) of 1.6 ml of thiopropionic acid (3 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition. The reaction was then left 18 hours at 50 °C.
Filtration of the reaction mixture and washing with 100 ml of distilled water gave the thiopropionic Sepharose derived gel.

D) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 151 µl of 3-mercapto-1,2-propanediol (0.3 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.
The reaction was left 1 hour under stirring at 20 °C. After 1 hour the reaction was treated with an aqueous solution (5ml dist.water) of 1.6 ml of thiopropionic acid (3 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition. The reaction was then left 18 hours at 50 °C.
Filtration of the reaction mixture and washing with 100 ml of distilled water gave the thiopropionic Sepharose derived gel.

Example 3.2. Anion exchange media

Gradients of immobilised trimethylamine and 3-mercapto-1,2-propanediol:

Bromine was added to a stirred suspension of 100 ml of allyl activated Sepharose™ 6 FF (0.3 mmol of allyl/ml), 4 g of AcONa and 100 ml of distilled water, till a persistent yellow colour was obtained. Sodium formate was then added till the suspension was fully decolourised.

The reaction mixture was filtered and the gel washed with 500 ml of distilled water. The activated gel was then directly transfer to 4 distinct reaction vessels:

A) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 251 µl of 3-mercapto-1,2-propanediol (0.5 equivalents per allyl group) and 2.4 g of NaCl, which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.

The reaction was left 1 hour under stirring at 50 °C. After 1 hour the reaction was treated with an aqueous solution (5ml dist.water) of 2.86 g of trimethylammonium chloride (5 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11.5 with 50 % aq. NaOH before the addition. The reaction was then left 18 hours at 50 °C.

The reaction was terminated by washing the gel on a glass filter funnel with 400 mL water, 80 ml 1M Sodium chloride and another 400 mL of water.

B) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 251 µl of 3-mercapto-1,2-propanediol (0.5 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.

The reaction was left 1 hour under stirring at 20 °C. After 1 hour the reaction was treated with an aqueous solution (5ml dist.water) of 2.86 g of trimethylammonium chloride (5 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11.5 with 50 % aq. NaOH before the addition. The reaction was then left 18 hours at 50 °C.

The reaction was terminated by washing the gel on a glass filter funnel with 400 mL water, 80 ml 1M Sodium chloride and another 400 mL of water.

C) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 151 µl of 3-mercapto-1,2-propanediol (0.3 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.
The reaction was left 1 hour under stirring at 50 °C. After 1 hour the reaction was treated with an aqueous solution (5ml dist.water) of 2.86 g of trimethylammonium chloride (5 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11.5 with 50 % aq. NaOH before the addition. The reaction was then left 18 hours at 50 °C.
The reaction was terminated by washing the gel on a glass filter funnel with 400 mL water, 80 ml 1M Sodium chloride and another 400 mL of water.

D) 20 ml of activated gel was treated with, an aqueous solution (10ml dist.water) of 151 µl of 3-mercapto-1,2-propanediol (0.3 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11,5 with 50 % aq. NaOH before the addition.
The reaction was left 1 hour under stirring at 20 °C. After 1 hour the reaction was treated with an aqueous solution (5ml dist.water) of 2.86 g of trimethylammonium chloride (5 equivalents per allyl group) and 2.4 g of NaCl which pH was adjusted to 11.5 with 50 % aq. NaOH before the addition. The reaction was then left 18 hours at 50 °C. The reaction was terminated by washing the gel on a glass filter funnel with 400 mL water, 80 ml 1M Sodium chloride and another 400 mL of water.

CLAIMS

1. A separation matrix comprised of ligands coupled to the surfaces of a porous support, wherein the ligands provide at least one chemical gradient in the support.
2. A separation matrix according to claim 1, wherein the support is comprised of at least one porous particle and the ligands provide at least one chemical gradient between the centre and the exterior surface of each porous particle.
3. A matrix according to claim 1 or 2, wherein at least one chemical gradient is a ligand density gradient resulting from a varying density of ligands present on the support.
4. A matrix according to claim 2 or 3, wherein the density gradient increases towards the centre of the support.
5. A matrix according to claim 2 or 3, wherein the density gradient decreases towards the centre of the support.
6. A matrix according to any one of the preceding claims, wherein at least one chemical gradient is the result of varying pKa values of functional groups of the ligands present on the support.
7. A matrix according to any one of the preceding claims, wherein at least one chemical gradient is the result of a varying net charge of the ligands present on the support.
8. A matrix according to any one of the preceding claims, wherein at least one gradient is a continuous and smooth gradient.
9. A matrix according to any one of the preceding claims, wherein the ligands present on the support provide at least two different functionalities.
10. A matrix according to claim 9, wherein said functionalities are selected from the group that consists of cation exchange ligands, anion exchange ligands, hydrophobic interaction chromatography (HIC) ligands, reversed phase chromatography (RPC) ligands, immobilised metal chelating ligands (IMAC), thiophilic ligands, and affinity ligands.
11. A matrix according to claim 9 or 10, wherein said at least two different functionalities are present on the same ligand.

12. A matrix according to claim 9, wherein the ligands present zwitterionic functionalities.

13. A matrix according to claim 9 or 10, wherein said at least two different functionalities are present on different ligand kinds, and each such ligand kind provides a separate chemical gradient within the support.

14. A chromatography column packed with a separation matrix comprised of ligands coupled to the surfaces of a porous support, wherein the ligands provide at least one chemical gradient within the support.

15. A chromatography column according to claim 14, wherein the support is comprised of at least one porous particle and the ligands provide at least one chemical gradient between the centre and the exterior surface of each porous particle.

16. A chromatography column according to claim 14 or 15, wherein at least one chemical gradient is a ligand density gradient.

17. A chromatography column according to any one of claims 14-16, which has been packed with a separation matrix according to any one of claims 1-13.

18. A method of preparing a separation matrix that comprises ligands coupled to the surfaces of a porous support, which method comprises the step of

(a) providing activatable groups on the surface of a porous support;

(b) activating said groups with an activation agent;

(c) reacting groups activated according to step b) with a compound which comprises at least one functionality;

wherein control of the reactivity in step (c) results in at least one chemical gradient within the support.

19. A method according to claim 18, wherein the reactivity is controlled by the concentration of the compound that comprises the functionalities in step (c).

20. A method according to claim 18-19, wherein the activatable groups of step (a) are carbon-carbon double bonds.

21. A method according to any one of claims 18-20, which also comprises to provide a the activatable groups present at the surface of at least one porous particle in a step preceding step (a).

22. A method according to claim 21, wherein the step preceding step (a) comprises to allylate hydroxyl groups present on the surface of a porous support.

23. A method according to claim 21 or 22, wherein steps (a)-(c) are replaced by a single step wherein an activated ligand is reacted with the activatable groups present on the surface of the support.

24. A method according to any one of claims 18-23, wherein the activation agent used in step (b) is a halogen.

25. A method according to any one of claims 18-24, wherein at least one chemical gradient is a ligand density gradient.

26. A method according to anyone of claims 18-24, wherein at least one chemical gradient is a continuous and smooth gradient.

27. A method according to any one of claims 18-26, wherein the at least two different functionalities are provided in step (c).

28. A method according to claim 27, wherein said at least two different functionalities are provided by one compound.

29. A matrix according to claim 27, wherein said at least two different functionalities are provided by different compounds.

30. A method according to any one of claims 18-29, wherein two or more chemical gradients are provided in the support, one of which is a ligand density gradient.

31. A method according to any one of claims 18-30, wherein the support is comprised of at least one porous particle, preferably an essentially spherical particle, and at least one gradient extends between the centre and the exterior surface of each porous particle.

32. A separation matrix prepared by the method according to any one of claims 18-31.

33. A process of liquid chromatography, wherein a liquid comprising at least one target molecule is contacted with a separation matrix that comprises ligands coupled to the surfaces of a porous support and the target molecule

is adsorbed to the matrix, wherein the ligands provide a chemical gradient within the support.

34. A process according to claim 33, which further comprises a step of eluting the adsorbed target molecule from the matrix by contacting the matrix with an eluent.

35. A process according to claim 33 or 34, wherein at least one chemical gradient is a ligand density gradient.

36. A process according to any one of claims 33-35, wherein the support is comprised of at least one porous particle and at least one chemical gradient extends between the centre and the exterior surface of each porous particle.

37. A process according to claim 33-36, wherein the functional groups of the ligands are zwitterions and the elution is performed at a pH different from that during the adsorption.

38. A process according to any one of claims 33-36, wherein the separation matrix is as defined in any one of claims 1-13.

ABSTRACT

The present invention relates to a separation matrix comprised of ligands coupled to the surfaces of a porous support, such as one or more porous particles, wherein the ligands provide at least one chemical gradient within the support. In the most advantageous embodiment, the chemical gradient is a ligand density gradient. The invention also relates to a method of preparing a separation matrix that comprises ligands coupled to the surfaces of a porous support, such as porous particles, which method comprises the steps of providing activatable groups, such as carbon-carbon double bonds, on the surface of a porous support; activating said groups with an activation agent, such as a halogen; and reacting the so activated groups with a compound which comprises at least one functionality. In the method according to the invention, the control of the diffusion rate in the second step results in at least one chemical gradient, such as a ligand density gradient, between the centre and the exterior surface of each porous particle.

Fig. 1

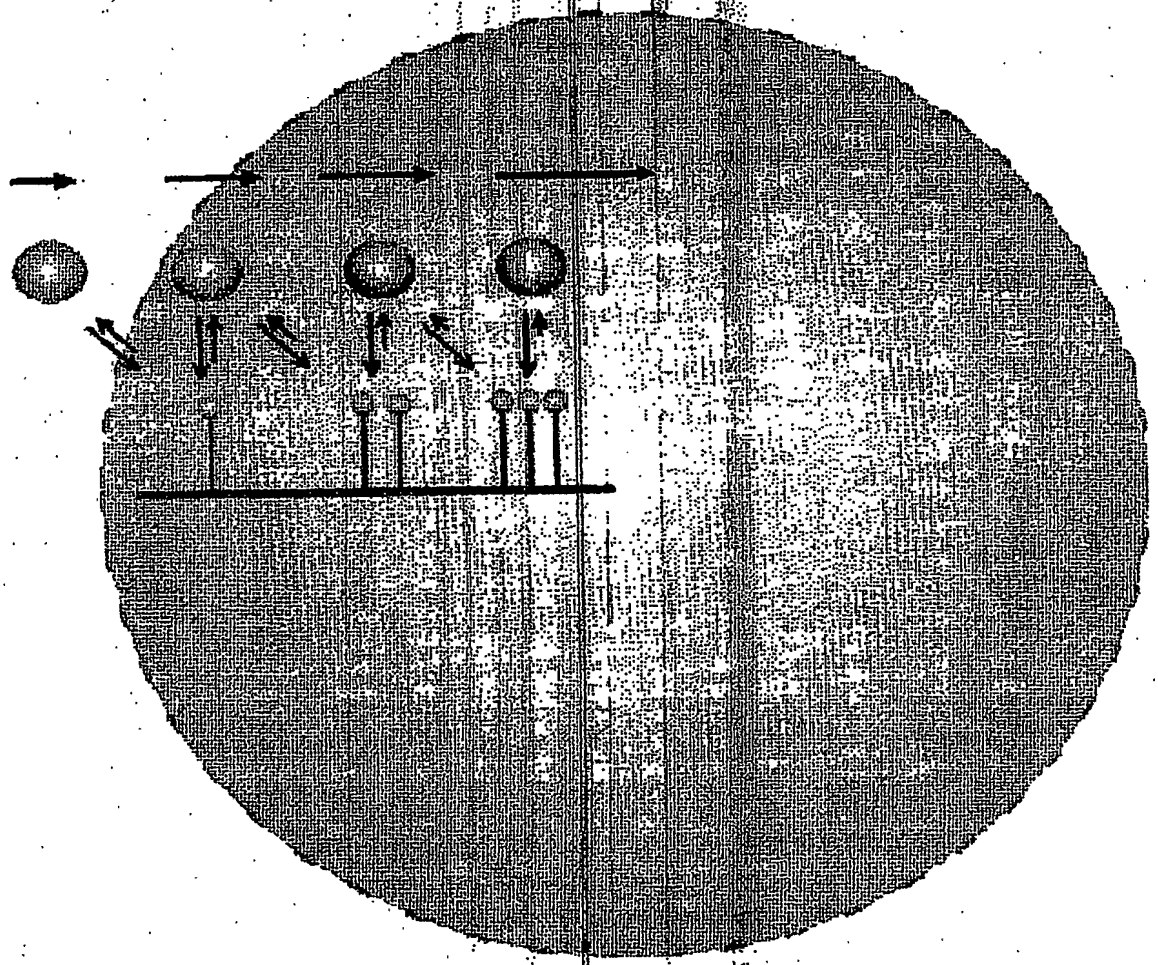

Fig. 2 A

Fig. 2 B

Fig. 2C

Fig. 2D

Fig. 2 E

Fig. 2 F

Fig. 2 G

Fig. 2 H